Amendments to the claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

1.- 42. (canceled)

43. (Currently Amended) A method of polarizing an immune response to an antigen in a subject, which method comprises administering to the subject a vaccine comprising one or more antigen(s) and an adjuvant composition comprising a Th1-activating alkaloid in an amount effective to polarize an immune response to the antigen(s) from type 2 towards type 1, wherein the alkaloid is selected from the group consisting of casuarine, 3, 7-diepi-casuarine, 7-epi-casuarine, 3, 6, 7-triepi-casuarine, 6,7-diepi-casuarine, 3-epi-casuarine, casuarine-6-α-D-glucoside, 3, 7-diepi-casuarine-6-α-glucoside, 7-epi-casuarine-6-α-D-glucoside, 3, 6, 7-triepi-casuarine-6-α-D-glucoside, 6, 7-diepi-casuarine-6-α-D-glucoside, and 3-epi-casuarine-6-α-D-glucoside.

has the formula:

$$RO \longrightarrow H \longrightarrow OH$$
 CH_2OH

wherein R is selected from the group comprising hydrogen, straight or branched, unsubstituted or substituted, saturated or unsaturated acyl, alkyl (e.g. cycloalkyl), alkenyl, alkynyl and aryl groups, or a pharmaceutically acceptable salt or acyl derivative thereof.

44. (Previously Presented) The method of claim 43 wherein the Th1-activating alkaloid stimulates the expression of IL-12 *in vitro* in lymphocytes and/or dendritic cells.

- 45. (Previously Presented) The method of claim 43 wherein the adjuvant composition further comprises an auxiliary adjuvant.
- 46. (Previously Presented) The method of claim 44 wherein the adjuvant composition further comprises an auxiliary adjuvant.
- 47. (Currently Amended) The method of claim 45 wherein the auxiliary adjuvant is selected from:
 - (a) a type 2 adjuvant; and/or
 - (b) a cytokine;
 - (c) a depot-forming agent;
 - (d) a saponin;
 - (e) a submicron oil-in-water emulsion;
 - (f) a CpG;
 - (g) a lipid A derivative;
 - (h) an MDP;
 - (i) an ISCOM®;
 - (j) an antigen-presenting cell (APC) (for example, a dendritic cell);
 - (k) a cytotoxic T lymphocyte (CTL); and
 - (1) a synergistic combination of any of the above.
- 48. (Currently Amended) The method of claim 46 wherein the auxiliary adjuvant is selected from:
 - (m)a type 2 adjuvant; and/or
 - (n) a cytokine;
 - (o) a depot-forming agent;
 - (p) a saponin;
 - (q) a submicron oil-in-water emulsion;
 - (r) a CpG;
 - (s) a lipid A derivative;
 - (t) an MDP;

- (u) an ISCOM®;
- (v) an antigen-presenting cell (APC) (for example, a dendritic cell);
- (w) a cytotoxic T lymphocyte (CTL); and
- a synergistic combination of any of the above.
- 49. (Previously Presented) The method of claim 43 wherein the vaccine is selected from: (a) a subunit vaccine; (b) a conjugate vaccine; (c) a DNA vaccine; (d) a recombinant vaccine; (e) a mucosal vaccine; (f) a therapeutic vaccine; (g) a prophylactic vaccine.
- 50. (Previously Presented) The method of claim 48 wherein the vaccine is selected from: (a) a subunit vaccine; (b) a conjugate vaccine; (c) a DNA vaccine; (d) a recombinant vaccine; (e) a mucosal vaccine; (f) a therapeutic vaccine; (g) a prophylactic vaccine.
- 51. (Currently Amended) The method of claim 43 wherein the one or more antigen(s) are selected from:
 - (a) nucleic acid(s) which encode one or more antigenic protein(s);
 - (b) protein(s) or peptide(s);
 - (c) glycoprotein(s);
 - (d) polysaccharide(s) (e.g. carbohydrate(s));
 - (e) fusion protein(s);
 - (f) lipid(s);
 - (g) glycolipid(s);
 - (h) peptide mimic(s) of polysaccharides;
 - (i) carbohydrate(s) and a protein(s) in admixture;
 - (j) carbohydrate-protein conjugate(s);
 - (k) cells or extracts thereof;
 - (1) dead or attenuated cells, or extracts thereof;
 - (m)tumour cells or extracts thereof;
 - (n) viral particles;
 - (o) allergen(s);
 - (p) mixtures of any of (a) to (o).

- 52. (Currently Amended) The method of claim 50 wherein the one or more antigen(s) are selected from:
 - (q) nucleic acid(s) which encode one or more antigenic protein(s);
 - (r) protein(s) or peptide(s);
 - (s) glycoprotein(s);
 - (t) polysaccharide(s) (e.g. carbohydrate(s));
 - (u) fusion protein(s);
 - (v) lipid(s);
 - (w)glycolipid(s);
 - (x) peptide mimic(s) of polysaccharides;
 - (y) carbohydrate(s) and a protein(s) in admixture;
 - (z) carbohydrate-protein conjugate(s);
 - (aa) cells or extracts thereof;
 - (bb) dead or attenuated cells, or extracts thereof;
 - (cc) tumour cells or extracts thereof;
 - (dd) viral particles;
 - (ee) allergen(s);
 - (ff) mixtures of any of (a) to (o).
- 53. (Previously Presented) The method of claim 51 wherein the one or more antigen(s) comprise a bacterial antigen, a viral antigen, a fungal antigen, a protozoal antigen, a prion antigen, a neoantigen, a tumour-associated antigen or a self-antigen.
- 54. (Previously Presented) The method of claim 52 wherein the one or more antigen(s) comprise a bacterial antigen, a viral antigen, a fungal antigen, a protozoal antigen, a prion antigen, a neoantigen, a tumour-associated antigen or a self-antigen.
- 55. (Previously Presented) The method of claim 51 wherein the one or more antigen(s) are dose-spared.

- 56. (Previously Presented) The method of claim 54 wherein the one or more antigen(s) are dose-spared.
- 57. (Previously Presented) The method of claim 43 wherein the vaccine is administered orally, mucosally, topically, epicutaneously, intramuscularly, intradermally, subcutaneously, intranasally, intravaginally, sublingually or *via* inhalation.
- 58. (Previously Presented) The method of claim 56 wherein the vaccine is administered orally, mucosally, topically, epicutaneously, intramuscularly, intradermally, subcutaneously, intranasally, intravaginally, sublingually or *via* inhalation.
- 59. (Canceled)
- 60. (Canceled)
- 61. (Canceled)
- 62. (Canceled)